

# Datasheet for ABIN6256503 anti-ERK1 antibody (pThr202, pTyr204)

4 Images

23 Pub

Publications



# Overview

| Quantity:            | 100 µL  |
|----------------------|---|
| Target:              | ERK1 (MAPK3)  |
| Binding Specificity: | pThr202, pTyr204  |
| Reactivity:          | Human, Mouse, Rat   |
| Host:                | Rabbit  |
| Clonality:           | Polyclonal  |
| Conjugate:           | This ERK1 antibody is un-conjugated   |
| Application:         | Western Blotting (WB), ELISA, Immunohistochemistry (IHC), Immunofluorescence (IF), Immunocytochemistry (ICC)                        |
| Product Details      |   |
| Immunogen:           | A synthesized peptide derived from human ERK1/2 around the phosphorylation site of Thr202/Tyr204.                                   |
| Isotype:             | lgG   |
| Specificity:         | Phospho-ERK1/2 (Thr202/Tyr204) Antibody detects endogenous levels of ERK1/2 only when phosphorylated at Threonine 202/Tyrosine 204. |

Predicted Reactivity:Pig,Zebrafish,Bovine,Horse,Sheep,RabbitPurification:The antibody is from purified rabbit serum by affinity purification via sequential<br/>chromatography on phospho- and non-phospho-peptide affinity columns.

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| Target:           | ERK1 (MAPK3)  |
|-------------------|---|
| Alternative Name: | MAPK3,MAPK1 (MAPK3 Products)  |
| Background:       | Description: Serine/threonine kinase which acts as an essential component of the MAP kinase         |
|                   | signal transduction pathway. MAPK1/ERK2 and MAPK3/ERK1 are the 2 MAPKs which play an                |
|                   | important role in the MAPK/ERK cascade. They participate also in a signaling cascade initiated      |
|                   | by activated KIT and KITLG/SCF. Depending on the cellular context, the MAPK/ERK cascade             |
|                   | mediates diverse biological functions such as cell growth, adhesion, survival and differentiation   |
|                   | through the regulation of transcription, translation, cytoskeletal rearrangements. The              |
|                   | MAPK/ERK cascade plays also a role in initiation and regulation of meiosis, mitosis, and            |
|                   | postmitotic functions in differentiated cells by phosphorylating a number of transcription          |
|                   | factors. About 160 substrates have already been discovered for ERKs. Many of these                  |
|                   | substrates are localized in the nucleus, and seem to participate in the regulation of transcription |
|                   | upon stimulation. However, other substrates are found in the cytosol as well as in other cellular   |
|                   | organelles, and those are responsible for processes such as translation, mitosis and apoptosis      |
|                   | Moreover, the MAPK/ERK cascade is also involved in the regulation of the endosomal                  |
|                   | dynamics, including lysosome processing and endosome cycling through the perinuclear                |
|                   | recycling compartment (PNRC), as well as in the fragmentation of the Golgi apparatus during         |
|                   | mitosis. The substrates include transcription factors (such as ATF2, BCL6, ELK1, ERF, FOS,          |
|                   | HSF4 or SPZ1), cytoskeletal elements (such as CANX, CTTN, GJA1, MAP2, MAPT, PXN,                    |
|                   | SORBS3 or STMN1), regulators of apoptosis (such as BAD, BTG2, CASP9, DAPK1, IER3, MCL1              |
|                   | or PPARG), regulators of translation (such as EIF4EBP1) and a variety of other signaling-related    |
|                   | molecules (like ARHGEF2, FRS2 or GRB10). Protein kinases (such as RAF1, RPS6KA1/RSK1,               |
|                   | RPS6KA3/RSK2, RPS6KA2/RSK3, RPS6KA6/RSK4, SYK, MKNK1/MNK1, MKNK2/MNK2,                              |
|                   | RPS6KA5/MSK1, RPS6KA4/MSK2, MAPKAPK3 or MAPKAPK5) and phosphatases (such as                         |
|                   | DUSP1, DUSP4, DUSP6 or DUSP16) are other substrates which enable the propagation the                |
|                   | MAPK/ERK signal to additional cytosolic and nuclear targets, thereby extending the specificity      |
|                   | of the cascade.   |
|                   | Gene: MAPK3   |
| Molecular Weight: | 42,44kDa  |
| Gene ID:          | 5595, 5594  |
| UniProt:          | P27361, P28482  |
| Pathways:         | MAPK Signaling, RTK Signaling, Interferon-gamma Pathway, Fc-epsilon Receptor Signaling              |
|                   | Pathway, Neurotrophin Signaling Pathway, Response to Growth Hormone Stimulus, Activation            |
|                   | of Innate immune Response, Cellular Response to Molecule of Bacterial Origin, Hepatitis C,          |

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# Application Details

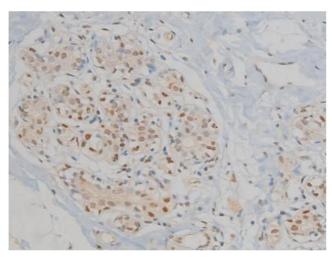
| Application Notes: | WB 1:500-1:2000, IHC 1:50-1:200, IF/ICC 1:200, ELISA(peptide) 1:20000-1:40000  |
|--------------------|--|
| Restrictions:      | For Research Use only  |
| Handling           |  |
| Format:            | Liquid   |
| Concentration:     | 1 mg/mL  |
| Buffer:            | Rabbit IgG in phosphate buffered saline , pH 7.4, 150 mM NaCl, 0.02 % sodium azide and 50 % glycerol.                  |
| Preservative:      | Sodium azide   |
| Precaution of Use: | This product contains Sodium azide: a POISONOUS AND HAZARDOUS SUBSTANCE which should be handled by trained staff only. |
| Storage:           | -20 °C   |
| Storage Comment:   | Store at -20 °C. Stable for 12 months from date of receipt.  |
| Expiry Date:       | 12 months  |
| Publications       |  |
| Product cited in:  | Deng, Cheng, Wu, Wang, Zhou, Huang: "Oxabicycloheptene Sulfonate Protects Against $\beta$ -                            |
|                    | Amyloid-induced Toxicity by Activation of PI3K/Akt and ERK Signaling Pathways Via GPER1 in                             |
|                    | C6 Cells." in: Neurochemical research, Vol. 42, Issue 8, pp. 2246-2256, (2018) (PubMed).                               |
|                    | Li, Xiong, Xu, Duan, Yang, Zhou, Tu: "miR-29a regulated ER-positive breast cancer cell growth                          |
|                    | and invasion and is involved in the insulin signaling pathway." in: Oncotarget, Vol. 8, Issue 20,                      |
|                    | pp. 32566-32575, (2018) (PubMed).  |
|                    | Xie, Cao, Yang, Xu, Wei, Wang: "Relaxin Attenuates Contrast-Induced Human Proximal Tubular                             |
|                    | Epithelial Cell Apoptosis by Activation of the PI3K/Akt Signaling Pathway In Vitro." in: BioMed                        |
|                    | research international, Vol. 2017, pp. 2869405, (2018) (PubMed).   |

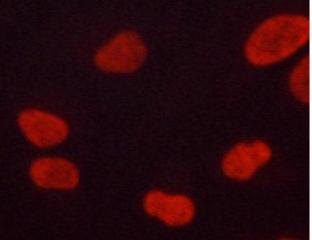
International: +49 (0)241 95 163 153 | USA & Canada: +1 877 302 8632 | support@antibodies-online.com Page 3/5 | Product datasheet for ABIN6256503 | 07/25/2024 | Copyright antibodies-online. All rights reserved. Peng, Wu, Deng, Zhou, Song, Yang, Zhang, Xu, Xia, Cai, Liu, Peng: "MiR-377 promotes white adipose tissue inflammation and decreases insulin sensitivity in obesity via suppression of sirtuin-1 (SIRT1)." in: **Oncotarget**, Vol. 8, Issue 41, pp. 70550-70563, (2018) (PubMed).

Li, Zhang, Jin, Zou, Wang, Hao, Fu, Jiao, Zhang, Lin, Matsuzaki, Zhao: "Dysifragilone A inhibits LPS-induced RAW264.7 macrophage activation by blocking the p38 MAPK signaling pathway." in: **Molecular medicine reports**, Vol. 17, Issue 1, pp. 674-682, (2018) (PubMed).

There are more publications referencing this product on: Product page

## Images





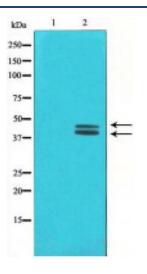
### Immunohistochemistry

**Image 1.** ABIN6267060 at 1/200 staining Human heart tissue sections by IHC-P. The tissue was formaldehyde fixed and a heat mediated antigen retrieval step in citrate buffer was performed. The tissue was then blocked and incubated with the antibody for 1.5 hours at 22°C. An HRP conjugated goat anti-rabbit antibody was used as the secondary.

#### Immunofluorescence (fixed cells)

**Image 2.** ABIN6267060 staining lovo cells by ICC/IF. Cells were fixed with PFA and permeabilized in 0.1% saponin prior to blocking in 10% serum for 45 minutes at 37°C. The primary antibody was diluted 1/400 and incubated with the sample for 1 hour at 37°C. A Alexa Fluor® 594 conjugated goat polyclonal to rabbit IgG (H+L), diluted 1/600 was used as secondary antibody.

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### Western Blotting

**Image 3.** Western blot analysis of ERK1/2 phosphorylation expression in HeLa whole cell lysates,The lane on the left is treated with the antigen-specific peptide.

Please check the product details page for more images. Overall 4 images are available for ABIN6256503.